Clean Version of Amended Claims

1 (Original). A compound of formula (I)

$$\begin{array}{c|c} X & Z & O \\ \hline & & \\ &$$

wherein

one of X and Y is silicon, and the other is carbon or silicon;

Z is oxygen, sulphur or -N(R)-, wherein R is hydrogen or alkyl;

R¹ is hydrogen, halogen, alkyl, alkenyl, alkynyl, alkoxy or cycloalkyl; and

R² is alkyl, alkenyl, alkynyl, cycloalkyl, heterocycloalkyl, aryl, heteroaryl,

-alkyl-cylcoalkyl, -alkyl-heterocycloalkyl, -alkyl-aryl or -alkyl-heteroaryl;

or a pharmaceutically acceptable salt thereof.

- 2 (Currently amended). The compound according to claim 1, wherein X and Y are each silicon.
- 3 (Currently amended). The compound according to claim 1, wherein \boldsymbol{Z} is oxygen.
- 4 (Currently amended). The compound according to claim 1, wherein R¹ is hydrogen or alkyl.
- 5 (Currently amended). The compound according to claim 4, wherein R¹ is methyl.
- 6 (Currently amended). The compound according to claim 1, wherein R^2 is aryl, -CH₂-cycloalkyl, CH₂-aryl, CH₂-heterocylcoaryl or CH₂-heteroaryl.
- 7 (Currently amended). The compound according to claim 6, wherein R^2 is optionally substituted phenyl.

8 (Currently amended). The compound according to claim 7, wherein R² is phenyl substituted with one, two or three alkoxy groups.

9 (Currently amended). The compound according to claim 1, which is 5-[(3,5,5,8,8-pentamethyl-5,8-disila-5,6,7,8-tetrahydro-2-naphthyl)methyl]-*N*-(2,4,6-trimethoxyphenyl)furan-2-carboxamide.

10 (Canceled).

11 (Currently amended). A pharmaceutical composition comprising a compound of formula (I)

$$X$$
 Z
 NH
 R^{1}
 R^{2}
 (I)

wherein

one of X and Y is silicon, and the other is carbon or silicon;

Z is oxygen, sulphur or -N(R)-, wherein R is hydrogen or alkyl;

R¹ is hydrogen, halogen, alkyl, alkenyl, alkynyl, alkoxy or cycloalkyl; and

R² is alkyl, alkenyl, alkynyl, cycloalkyl, heterocycloalkyl, aryl, heteroaryl,

-alkyl-cylcoalkyl, -alkyl-heterocycloalkyl, -alkyl-aryl or -alkyl-heteroaryl;

or a pharmaceutically acceptable salt thereof; and a pharmaceutically acceptable diluent or carrier.

12 (Currently amended). A method_for the treatment or prevention of a disease or condition associated with GnRH wherein said method comprises administering a compound of formula (I)

$$\begin{array}{c|c} X & Z & O \\ & & NH \\ & & R^2 \end{array}$$
 (I)

wherein

one of X and Y is silicon, and the other is carbon or silicon;

Z is oxygen, sulphur or -N(R)-, wherein R is hydrogen or alkyl;

R1 is hydrogen, halogen, alkyl, alkenyl, alkoxy or cycloalkyl; and

R² is alkyl, alkenyl, alkynyl, cycloalkyl, heterocycloalkyl, aryl, heteroaryl,

-alkyl-cylcoalkyl, -alkyl-heterocycloalkyl, -alkyl-aryl or -alkyl-heteroaryl;

or a pharmaceutically acceptable salt thereof;

to a patient in need of such treatment.

- 13 (Currently amended). The method according to claim 12, for the treatment or prevention of progression of cancer.
- 14 (Currently amended). The method according to claim 13, for leukaemia therapy.
- 15 (Currently amended). The method according to claim 12, for the treatment or prevention of a fertility disorder.
- 16 (Currently amended). The method according to claim 12, for the treatment or prevention of HIV infection or AIDS.
- 17 (Currently amended). The method according to claim 12, for the treatment or prevention of Alzheimer's disease.
- 18 (Currently amended). The method according to claim 12, for the treatment or prevention of fibrosis.

- 19 (Currently amended). The method according to claim 12, for the treatment or prevention of endometriosis.
- 20 (Currently amended). The method according to claim 12, for the treatment or prevention of uterine fibroids.
- 21(Currently amended). The method according to claim 12, for the treatment or prevention of uterine leiomyoma.